IN THE CLAIMS

- 1. (Currently Amended) Non-human <u>transgenic</u> animal, being [transgenic] <u>transgenic</u> for
- an <u>anti-NGH (Nerve Growth Factor)</u> antibody [or fragments thereof and] having a phenotype
- 3 reminiscent of a human [pathology] neurodegenerative syndromes, muscular atrophy or
- 4 <u>dystrophy or immune disorders</u>.
- 1 2. (Cancelled).
- 3. (Currently Amended) A non-human transgenic animal according to claim [2] 1
- wherein the [human pathology is the] phenotype recapitulates the features of the human
- 3 Alzheimer disease (AD).
- 4. (Currently Amended) A non-human transgenic animal according to claim 3 exhibiting
- 2 at least one of the anatomical, histological, molecular or phenotypic markers included in the
- 3 following group: deposition in Central Nervous System (CNS) of plaques of amyloid precursor
- 4 protein (APP) or of β-amyloid protein, hyperphosphorylation of the tau protein, neurofibrillar
- 5 pathology, and deficits in the cholinergic system.
- 5. (Currently Amended) A non-human transgenic animal according to claim 4 further
- 2 exhibiting at least one of the anatomical, histological, molecular or phenotypic markers included
- in the following group: glial activation, neuronal loss, cortical and hippocampal atrophy, and
- 4 muscular myositis.
- 6. (Currently Amended) A non-human transgenic animal according to claim 5 exhibiting
- the following anatomical, histological, molecular or phenotypic markers: deposition in Central
- Nervous System (CNS) of plaques of amyloid precursor protein (APP) or of β-amyloid protein,

- 4 hyperphosphorylation of the tau protein, neurofibrillar pathology, deficits in the cholinergic
- system, glial activation, neuronal loss, cortical and hippocampal atrophy, and muscular myositis.
- 7. (Currently Amended) A non-human transgenic animal according to claim 6 exhibiting
- 2 the anatomical, histological, molecular or phenotypic markers as defined [in-Table 1] by
- 3 decrease of cortical thickness, hippocampal formation atrophia, ventricle dilation, cognitive
- 4 <u>deficits, neuronal loss, apoptosis, β-amyloid plaques, hyperphosphorylated tau, neurofibrillary</u>
- 5 tangles, tau aggregates, dystrophic neuritis, glial activation, cholinergic deficit, synaptic loss,
- 6 decreased synaptic plasticity, skeletal muscle atrophia and dystrophy, amyloid deposits in
- 7 skeletal muscles, hyperphosphorylated tau in skeletal muscles, inflammation in skeletal muscles,
- 8 vacuolization of myofibers, increased number of central nuclei in myofibers, or spleen
- 9 alterations.

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- 8. (Original) A non-human transgenic animal according to claim 7 wherein said markers
- 2 are expressed in the adult age.
- 9. (Original) A non-human transgenic animal according to claim 7 wherein the
- 2 occurrence of the tau hyperphosphorylation and/or the β-amyloid protein deposition in the back
- 3 or lower limb skeletal muscles and/or the atrophy of said skeletal muscles are present
- 4 concomitantly to the earliest occurrence of other neurological markers.
 - 10. (Cancelled)
- 1 11. (Original) A non-human transgenic animal according to claim 1 wherein the anti-
- 2 NGF antibody blocks the binding of NGF to its receptors.

- 1 12. (Original) A non-human transgenic animal according to claim 1 wherein the anti-
- 2 NGF antibody is expressed mainly in adulthood
- 1 13. (Original) A non-human transgenic animal according to claim 12 wherein the anti-
- 2 NGF antibody levels in the serum of the adult animal are comprised between 50 ng/ml and 500
- 3 ng/ml.
- 1 14. (Currently Amended) A non-human transgenic animal according to claim [10] 1
- wherein the anti-NGF antibody is the monoclonal anti-NGF α D11 antibody.
- 15. (Original) A non-human transgenic animal according to claim 14 wherein the αD11
- 2 antibody is a α D11 chimeric antibody.
- 1 16. (Original) A non-human transgenic animal according to claim 15 wherein the
- 2 chimeric antibody is a humanised chimeric antibody.
- 1 17. (Previously Presented) A non-human transgenic animal according to claim 1, wherein
- the animal is a mammalian.
- 1 18. (Original) A non-human transgenic animal according to claim 17 belonging to the
- 2 murine genus.
- 1 19. A non-human transgenic animal according to claim 18 belonging to the Mus musculus
- 2 [BS6JL] B6SJL strain.
- 1 20-37.(Cancelled)